In the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1-26. (Canceled)

27. (Currently amended) A **compound** peptide which binds to a DM2 protein, which **compound** peptide comprises an amino acid motif comprising at least the eight consecutive amino acids from F to R₄ of the formula

$$R_1-X-F-X-R_2-R_3-W-X-X-R_4$$

(I) (SEQ ID NO: 4)

wherein

R₁ is a proline (P), leucine (L), glutamic acid (E), cysteine (C) or glutamine (Q),

X stands for any natural amino acid,

R₂ is arginine (R), histidine (H), glutamic acid (E), cysteine (C), serine (S), or aspartic acid (D),

R₃ is histidine (H), phenylalanine (F) or tyrosine (Y),

R₄ is phenylalanine (F), glutamine (Q) or leucine (L); and

F is phenylalanine and W is tryptophan,

and inhibits the binding of said DM2 protein to a p53 protein.

- 28. (Currently amended) The compound peptide according to claim 27 wherein the compound peptide binds to human DM2 (HDM2).
- 29. (Currently amended) The **compound peptide** according to claim 27, further comprising which is coupled to a biotin moiety coupled to the amino acid motif.
- 30. (Currently amended) The compound peptide according to claim 27, which is wherein the amino acid motif comprises a cyclic peptide.
- 31. (Currently amended) The **compound peptide** according to claim 27, **wherein the amino acid motif comprises which is** a cyclic lactam.
- 32. (Currently amended) The eompound peptide according to claim 27 wherein the amino acid motif which comprises a disulfide bond.

- 33. (Currently amended) The **eompound peptide** according to claim 27 which comprises no more than fifteen amino acids (15 mers).
- 34. (Currently amended) The **compound peptide** according to claim 27 which comprises an amino acid motif selected from the group consisting of M-P-R-F-M-D-Y-W-E-G-L-N (SEQ ID NO: 6), Q-P-T-F-S-D-Y-W-K-L-L-P (SEQ ID NO: 7), and P-X-F-X-D-Y-W-X-X-L (SEQ ID NO: 8).
- 35. (Currently amended) The compound according to claim 27, A peptide which comprises eight amino acids according to the formula

wherein R2 is arginine (R), histidine (H), glutamic acid (E), cysteine (C), serine (S), or aspartic acid (D);

R3 is histidine (H), phenylalanine (F), or tyrosine (Y);

R4 is phenylalanine (F), gutamine glutamine (Q) or leucine (L);

X2 is methionine (M), isoleucine (I), threonine (T), arginine (R), alanine (A) or serine (S);

X3 is glutamic acid (E), threonine (T), alanine (A), phenylalanine (F) or serine (S); and X4 is glycine (G), glutamine (Q), threonine (T), alanine (A) or aspartic acid (D).

36. (Currently amended) The **compound peptide** according to claim **27 35** comprising an amino acid motif of the formula

wherein

R2 is arginine (R), histidine (H), glutamic acid (E), eystine cysteine (C), serine (S), or aspartic acid (D);

R3 is histidine (H), phenylalanine (F) or tyrosine (Y);

R4 is phenylalanine (F), glutamine (Q) or leucine (L);

X1 is arginine (R), asparagine (N), alanine (A), threonine (T), or valine (V);

X2 is methionine (M), isoleucine (I), threonine (T), arginine (R), alanine (A), or serine (S);

X3 is glutamic acid (E), threonine (T), alanine (A), phenylalanine (F), or serine (S); and X4 is glycine (G), glutamine (Q), threonine (T), alanine (A), or aspartic acid (D).

- 37. (Canceled)
- 38. (Currently amended) The **eompound peptide** according to claim 27, wherein R2 is aspartic acid (D).
- 39. (Currently amended) The **compound** peptide according to claim 35, wherein at least one of R2, X2, X3, and X4 is defined as follows: R2 is aspartic acid (D), X2 is methionine (M), X3 is glutamic acid (E), and X4 is glycine (G).
- 40. (Currently amended) The **compound peptide** according to claim 36, wherein at least one of R2, X1, X2, X3, and X4 is defined as follows: R2 is aspartic acid (D), X1 is arginine (R), X2 is methionine (M), X3 is glutamic acid (E), and X4 is glycine (G).
- 41. (Currently amended) A method for inhibiting the binding of a DM2 protein to a p53 protein comprising contacting said DM2 protein with a **compound peptide** which **compound peptide** comprises an amino acid motif comprising at least eight consecutive amino acids of the formula

$$R_1-X-F-X-R_2-R_3-W-X-X-R_4$$
 (I) (SEQ ID NO: 4)

wherein

R₁ is a proline (P), leucine (L), glutamic acid (E), cysteine (C) or glutamine (Q),

X stands for any natural amino acid,

R₂ is arginine (R), histidine (H), glutamic acid (E), cysteine (C), serine (S), or aspartic acid (D),

R₃ is histidine (H), phenylalanine (F) or tyrosine (Y),

R₄ is phenylalanine (F), glutamine (Q) or leucine (L); and

F is phenylalanine and W is tryptophan,

and inhibits the binding of said DM2 protein to a p53 protein.

42. (Previously presented) The method of claim 41 wherein R2 is aspartic acid (D).

43-51. (Canceled)

52. (Currently amended) A composition comprising a **compound peptide**, which **compound peptide** comprises an amino acid motif comprising at least eight consecutive amino acids of the formula

$$R_1-X-F-X-R_2-R_3-W-X-X-R_4$$
 (I) (SEQ ID NO: 4)

wherein

R₁ is a proline (P), leucine (L), glutamic acid (E), cysteine (C) or glutamine (Q),

X stands for any natural amino acid,

R₂ is arginine (R), histidine (H), glutamic acid (E), cysteine (C), serine (S), or aspartic acid (D),

R₃ is histidine (H), phenylalanine (F) or tyrosine (Y),

R₄ is phenylalanine (F), glutamine (Q) or leucine (L); and

F is phenylalanine and W is tryptophan,

and inhibits the binding of said DM2 protein to a p53 protein, in admixture with at least one pharmaceutically acceptable carrier.